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(54) Title: **DIAPER HAVING A LOTIONED TOPSHEET CONTAINING A POLYSILOXANE EMOLIENT**

(57) Abstract

A diaper containing a liquid pervious topsheet coated with a lotion composition is disclosed. The lotion composition reduces the adherence of BM to the skin of the wearer, thereby improving the ease of BM cleanup. The lotion composition comprises a plastic or fluid polysiloxane emollient such as phenyl-functional polymethylsiloxanes, an immobilizing agent such as a fatty alcohol or paraffin wax to immobilize the emollient on the surface of the diaper topsheet, and optionally a hydrophilic surfactant to improve wettability of the coated topsheet. Because the polysiloxane emollient is substantially immobilized on the surface of the topsheet, less lotion is required to impart the desired therapeutic or protective lotion coating benefits.

the wearer. This procedure usually involves the caregiver's pouring of the oil or lotion, for example, in one of their hands, rubbing both hands together to distribute the substance thereon and then wiping the same on the skin of the infant. To eliminate the need for this wasteful, messy, and easily forgotten procedure, there have been numerous previous attempts to prepare absorbent articles which contain a protective or therapeutic skin care substance on the topsheet.

One substance that has been applied as a lotion to diaper products to impart a soothing, protective coating is mineral oil. Mineral oil (also known as liquid petrolatum) is a mixture of various liquid hydrocarbons obtained by distilling the high-boiling (i.e., 300°-390°C) fractions in petroleum. Mineral oil is liquid at ambient temperatures, e.g. 20°-25°C. As a result, mineral oil is relatively fluid and mobile, even when applied to diaper topsheets.

Because mineral oil is fluid and mobile at ambient temperatures, it tends not to remain localized on the surface of the topsheet, but instead migrates through the topsheet into the interior of the diaper. Accordingly, relatively high levels of mineral oil need to be applied to the topsheet to provide the desired therapeutic or protective coating lotion benefits. This leads not only to increased costs for these lotioned diaper products, but other detrimental effects as well.

One of these detrimental effects is a decrease in the fluid handling properties as high levels of mineral oil tend to block the topsheet openings. Also, as mineral oil migrates to the interior of the diaper, it tends to act as a hydrophobic additive, thus decreasing the absorbency of the underlying absorbent core, if one is used. This decrease in absorbency becomes more pronounced as the level of mineral oil applied is increased.

Even without increasing its level, the tendency of mineral oil to migrate once applied has other detrimental effects. For example, the applied mineral oil can transfer to, into and through the packaging or wrapper material for the lotioned diaper product. This can create the need for barrier-type packaging or wrapper films to avoid smearing or other leakage of mineral oil from the diaper product.

Accordingly, it would be desirable to provide lotioned diaper products that: (1) have desirable therapeutic or protective coating lotion benefits (2)

do not require relatively high levels of coatings that are liquid at room temperature (e.g., mineral oil) (3) do not adversely affect the absorbency of the diaper product; and (4) do not require special wrapping or barrier materials for packaging.

5 U.S. Patent 3,585,998 to Hayford et al. teaches a disposable baby diaper, an interior liner of which carries an array of pressure-rupturable capsules containing baby oil. The patent teaches that it is desirable to break the capsules prior to using the diaper by applying pressure with such household items as a rolling pin, hand iron, etc. The same principle of 10 pressure-rupturable capsules is used in U.S. Patent 3,464,413 to Goldfarb et al. for making bandages capable of delivering a medicinal material to an injury. Articles disclosed by both patents have a serious drawback. Namely, unless the capsules are ruptured by applying pressure prior to 15 using the diaper or the bandage, the skin-care substance contained in the capsules is either not delivered at all or is delivered non-uniformly leaving some areas of skin uncoated.

U.S. Patent 3,896,807 to Buchalter teaches an article impregnated with a solid oil phase of cream formulation which forms a cream upon addition of moisture thereto. A major disadvantage of the article disclosed 20 by the reference is that transfer of a beneficial substance from the absorbent substrate to skin is delayed and is only realized when body fluids are released.

U.S. Patent 3,489,148 to Duncan et al. teaches a baby diaper comprising a hydrophobic and oleophobic topsheet wherein a portion of the 25 topsheet is coated with a discontinuous film of oleaginous material. A major disadvantage of the diapers disclosed in the Duncan et al. reference is that the hydrophobic and oleophobic topsheets are slow in promoting transfer of urine to the underlying absorbent cores.

Therefore, it is an object of the present invention to provide a 30 disposable diaper having a hydrophilic topsheet having superior fluid handling properties.

It is a further object of the present invention to provide a hydrophilic diaper topsheet having a lotion coating on the outer surface of the topsheet that is transferable to the wearer's skin and is effective at reducing the 35 adherence of BM to the skin, thereby improving the use of BM cleanup.

It is yet a further object of the present invention desirable to provide lotioned diaper products that: (1) have desirable BM release, cleaning, therapeutic or protective lotion coating benefits (2) do not require relatively high levels of mineral oil; (3) do not adversely affect the fluid handling properties of the diaper; and (4) do not require special wrapping or barrier materials for packaging.

These and other objects are obtained using the present invention, as will become readily apparent from a reading of the following disclosure.

10

SUMMARY OF THE INVENTION

The present invention relates to a disposable diaper having a lotion coating on the outer surface of the topsheet that is semisolid or solid at ambient temperatures (i.e., at 20°C) and is adapted to be transferred to the wearer's skin, where it acts to reduce the adherence of BM to the skin of the wearer, thereby improving the ease of BM clean up.

Briefly, the disposable diapers of the present invention comprise:

- A) a liquid impervious backsheet;
- B) a liquid pervious, hydrophilic topsheet joined to said backsheet, said topsheet having an inner surface oriented toward the interior of said diaper and an outer surface oriented toward the skin of the wearer when said diaper is being worn, wherein at least a portion of said topsheet outer surface comprises an effective amount of a lotion coating which is semi-solid or solid at 20°C and which is partially transferable to the wearer's skin, said lotion coating comprising:
 - (i) from about 10 to about 95% of a substantially water free emollient having a plastic or fluid consistency at 20°C and comprising a member selected from the group consisting of petroleum-based emollients, fatty acid ester emollients, alkyl ethoxylate emollients, polysiloxane emollients, and mixtures thereof;

5 (ii) from about 5 to about 90% of an agent capable of immobilizing said emollient on said outer surface of the topsheet, said immobilizing agent having a melting point of at least about 35°C; and

5 C) an absorbent core positioned between said topsheet and said backsheet.

10 The quantity of lotion coating on at least a portion of the diaper topsheets of the present invention preferably ranges from about 0.1 mg/in² to about 25 mg/in², more preferably from about 1 mg/in² to about 10 mg/in². Lotioned diaper topsheets according to the present invention provide desirable BM cleaning, therapeutic or protective lotion coating benefits. Because the emollient is substantially immobilized on the surface of the topsheet, less lotion composition is needed to impart the desired skin care benefits. In addition, special barrier or wrapping materials are unnecessary in packaging the lotioned diaper products of the present invention.

20 As will be discussed hereinafter, the lotion compositions of the present invention preferably have a melting profile such that they are relatively immobile and localized on the diaper topsheet at room temperature, are transferable to the wearer at body temperature, and yet are not completely liquid under extreme storage conditions.

25 Importantly, the lotion compositions of the present invention are easily transferable to the skin by way of normal contact, wearer motion, and/or body heat. Without being bound by theory, it is believed that the lotion composition changes the surface energy of the skin, and/or forms a "barrier" reducing the affinity of the skin for BM. The BM, therefore, has a reduced tendency to stick to the skin, and is easier to remove.

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BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is a schematic representation illustrating a preferred process for applying the lotion composition of the present invention to diaper topsheets.

Figure 2 is a schematic representation illustrating an alternative process for applying the lotion composition of the present invention to diaper topsheets.

5 Figure 3 is an absorbent article in the form of a diaper according to the present invention.

DETAILED DESCRIPTION OF THE INVENTION

10 As used herein, the term "comprising" means that the various components, ingredients, or steps, can be conjointly employed in practicing the present invention. Accordingly, the term "comprising" encompasses the more restrictive terms "consisting essentially of" and "consisting of."

All percentages, ratios and proportions used herein are by weight unless otherwise specified.

15 A. Absorbent Article

20 As used herein, the term "absorbent article" refers to devices which absorb and contain body exudates, and more specifically, refers to devices which are placed against the skin of a wearer to absorb and contain the various exudates discharged from the body. The term "disposable" is used herein to describe absorbent articles which are not intended to be laundered or otherwise restored or reused as an absorbent article after a single use. Examples of disposable absorbent articles include feminine hygiene garments such as sanitary napkins and panti-liners, diapers, 25 incontinence briefs, diaper holders, training pants, and the like.

30 Disposable absorbent articles typically comprise a liquid pervious topsheet, a liquid impervious backsheet joined to the topsheet and an absorbent core positioned between the topsheet and the backsheet. Disposable absorbent articles and components thereof, including the topsheet, backsheet, absorbent core, and any individual layers of these components, have a body surface and a garment surface. As used herein, "body surface" means that surface of the article or component which is intended to be worn toward or adjacent to the body of the wearer, while the "garment surface" is on the opposite side and is intended to be worn toward

or placed adjacent to the wearer's body or undergarments when the disposable absorbent article is worn.

5 The following description generally discusses the absorbent core, topsheet, and backsheet materials that are useful in disposable absorbent articles. It is to be understood that this general description applies to these components of the specific absorbent articles shown in Figure 3 and further described below, in addition to those of other disposable absorbent articles which are generally described herein.

10 In general, the absorbent core is capable of absorbing or retaining liquids (e.g., menses, urine, and/or other body exudates). The absorbent core is preferably compressible, conformable, and non-irritating to the wearer's skin. The absorbent core may be manufactured in a wide variety of sizes and shapes (e.g., rectangular, oval, hourglass, "T" shaped, dog bone, asymmetric, etc.). In addition to the absorbent composites of the 15 present invention, the absorbent core may include any of a wide variety of liquid-absorbent materials commonly used in absorbent articles, such as comminuted wood pulp, which is generally referred to as airfelt. Examples of other suitable absorbent materials for use in the absorbent core include creped cellulose wadding; meltblown polymers including coform; chemically 20 stiffened, modified or cross-linked cellulosic fibers; synthetic fibers such as crimped polyester fibers; peat moss; tissue including tissue wraps and tissue laminates; absorbent foams; absorbent sponges; superabsorbent polymers; absorbent gelling materials; or any equivalent material or combinations of materials, or mixtures of these.

25 The configuration and construction of the absorbent core may also be varied (e.g., the absorbent core may have varying caliper zones and/or have a profile so as to be thicker in the center; hydrophilic gradients; gradients of the absorbent composite of the present invention, superabsorbent gradients; or lower average density and lower average 30 basis weight zones, e.g., acquisition zones; or may comprise one or more layers or structures). The total absorbent capacity of the absorbent core should, however, be compatible with the design loading and the intended use of the absorbent article. Further, the size and absorbent capacity of the absorbent core may be varied to accommodate different uses such as 35 diapers, incontinence pads, pantliners, regular sanitary napkins, and

overnight sanitary napkins, and to accommodate wearers ranging from infants to adults.

5 The absorbent core can include other absorbent components that are often used in absorbent articles, for example, a dusting layer, a wicking or acquisition layer, or a secondary topsheet for increasing the wearer's comfort.

10 The topsheet is preferably compliant, soft feeling, and non-irritating to the wearer's skin. Further, the topsheet is liquid pervious, permitting liquids (e.g., menses and/or urine) to readily penetrate through its thickness. A suitable topsheet may be manufactured from a wide range of materials such as woven and nonwoven materials (e.g., a nonwoven web of fibers); polymeric materials such as apertured formed thermoplastic films, apertured plastic films, and hydroformed thermoplastic films; porous foams; reticulated foams; reticulated thermoplastic films; and thermoplastic scrims. 15 Suitable woven and nonwoven materials can be comprised of natural fibers (e.g., wood or cotton fibers), synthetic fibers (e.g., polymeric fibers such as polyester, polypropylene, or polyethylene fibers) or from a combination of natural and synthetic fibers. When the topsheet comprises a nonwoven web, the web may be manufactured by a wide number of known techniques. 20 For example, the web may be spunbonded, carded, wet-laid, melt-blown, hydroentangled, combinations of the above, or the like.

25 The backsheet is impervious to liquids (e.g., menses and/or urine) and is preferably manufactured from a thin plastic film, although other flexible liquid impervious materials may also be used. As used herein, the term "flexible" refers to materials which are compliant and will readily conform to the general shape and contours of the human body. The backsheet prevents the exudates absorbed and contained in the absorbent core from wetting articles which contact the absorbent article such as bedsheets, pants, pajamas and undergarments. The backsheet may thus 30 comprise a woven or nonwoven material, polymeric films such as thermoplastic films of polyethylene or polypropylene, or composite materials such as a film-coated nonwoven material. A suitable backsheet is a polyethylene film having a thickness of from about 0.012 mm (0.5 mil) to about 0.051 mm (2.0 mils). Exemplary polyethylene films are manufactured by Clopay Corporation of Cincinnati, Ohio, under the designation P18-1401 35 and by Tredegar Film Products of Terre Haute, Indiana, under the

5 designation XP-39385. The backsheet is preferably embossed and/or matte finished to provide a more clothlike appearance. Further, the backsheet may permit vapors to escape from the absorbent core (i.e., the backsheet is breathable) while still preventing exudates from passing through the backsheet. The size of the backsheet is dictated by the size of the absorbent core and the exact absorbent article design selected.

10 The backsheet and the topsheet are positioned adjacent the garment surface and the body surface, respectively, of the absorbent core. The absorbent core is preferably joined with the topsheet, the backsheet, or both in any manner as is known by attachment means (not shown in Figure 3) such as those well known in the art. However, embodiments of the present invention are envisioned wherein portions of the entire absorbent core are unattached to either the topsheet, the backsheet, or both.

15 For example, the backsheet and/or the topsheet may be secured to the absorbent core or to each other by a uniform continuous layer of adhesive, a patterned layer of adhesive, or an array of separate lines, spirals, or spots of adhesive. Adhesives which have been found to be satisfactory are manufactured by H. B. Fuller Company of St. Paul, Minnesota under the designation HL-1258 or H-2031. The attachment means will preferably comprise an open pattern network of filaments of adhesive as is disclosed in U.S. Patent 4,573,986, issued to Minetola, et al. on March 4, 1986, and which is incorporated herein by reference. An exemplary attachment means of an open pattern network of filaments comprises several lines of adhesive filaments swirled into a spiral pattern 20 such as illustrated by the apparatus and method shown in U.S. Patent 3,911,173 issued to Sprague, Jr. on October 7, 1975; U.S. Patent 4,785,996 issued to Zwieker, et al. on November 22, 1978; and U.S. Patent 4,842,666 issued to Weranicz on June 27, 1989. Each of these patents are incorporated herein by reference. Alternatively, the attachment means may 25 comprise heat bonds, pressure bonds, ultrasonic bonds, dynamic mechanical bonds, or any other suitable attachment means or combinations of these attachment means as are known in the art.

30 A preferred disposable absorbent article in which the lotioned topsheets of the present invention may be used are diapers. As used herein, the term "diaper" refers to an absorbent article generally worn by infants, and incontinent persons that is worn about the low torso of the

wearer. In other words, the term "diaper" includes infant diapers, training pants, adult incontinence devices, etc.

Figure 3 is a plan-view of the diaper 50 of the present invention in its flat-out, uncontracted state (i.e., with elastic induced contraction pulled out) with portions of the structure being cut-away to more clearly show the construction of the diaper 50 and with the portion of the diaper 50 which faces away from the wearer, the outer surface, oriented towards the viewer. As shown in Figure 3, the diaper 50 preferably comprises a liquid pervious topsheet 520; a liquid impervious backsheet 530 joined with the topsheet 520; an absorbent core 540 positioned between the topsheet 520 and the backsheet 530, the absorbent core 540 having a garment facing surface 542, a body facing surface 544, side edges 546, waist edges 548, and ears 549. The diaper 50 preferably further comprises elasticized leg cuffs 550; an elastic waist feature multiply designated as 560; and a fastening system generally multiply designated as 570.

The diaper 50 is shown in Figure 3 to have an outer surface 52, an inner surface 54 opposed to the outer surface 52, a first waist region 56, a second waist region 58, and a periphery 51 which is defined by the outer edges of the diaper 50 in which the longitudinal edges are designated 55 and the end edges are designated 57. (While the skilled artisan will recognize that a diaper is usually described in terms of having a pair of waist regions and a crotch region between the waist regions, in this application, for simplicity of terminology, the diaper 50 is described as having only waist regions including a portion of the diaper which would typically be designated as part of the crotch region). The inner surface 54 of the diaper 50 comprises that portion of the diaper 50 which is positioned adjacent to the wearer's body during use (i.e., the inner surface 54 generally is formed by at least a portion of the topsheet 520 and other components that may be joined to the topsheet 520). The outer surface 52 comprises that portion of the diaper 50 which is positioned away from the wearer's body (i.e., the outer surface 52 generally is formed by at least a portion of the backsheet 530 and other components that may be joined to the backsheet 530). (As used herein, the portion of the diaper 50 or component thereof which faces the wearer is also referred to as the body facing surface. Similarly, the portion facing away from the wearer is also referred to herein as the garment facing surface.) The first waist region 56 and the second waist region 58 extend,

respectively, from the end edges 57 of the periphery 51 to the lateral centerline 53 of the diaper 50. Figure 3 also shows the longitudinal centerline 59.

Figure 3 shows a preferred embodiment of the diaper 50 in which the 5 topsheet 520 and the backsheet 530 have length and width dimensions generally larger than those of the absorbent core 540. The elasticized leg cuffs 550 and the backsheet 530 extend beyond the edges of the absorbent core 540 to thereby form the periphery 51 of the diaper 50.

10 Diapers of the present invention can have a number of well known configurations, with the absorbent cores thereof being adapted to the present invention. Exemplary configurations are described generally in U.S. Patent 3,860,003 issued to Buell on January 14, 1975; U.S. Patent 5,151,092 issued to Buell et al. on September 29, 1992; U.S. Patent 5,221,274 issued to Buell et al. on June 22, 1993. Each of these patents is 15 incorporated herein by reference. Another diaper configuration to which the present invention can be readily adapted are described in co-pending U.S. Patent Application Serial Nos. 08/203,456; filed on February 28, 1994 and incorporated herein by reference. The absorbent cores of diapers described in these patents can be adapted in light of the teachings herein 20 to include the absorbent composite of the present invention as an absorbent gelling material described therein.

A topsheet 520 which is particularly suitable for use in the diaper 50, is carded and thermally bonded by means well known to those skilled in the fabrics art. A satisfactory topsheet for the present invention comprises 25 staple length polypropylene fibers having a denier of about 2.2. As used herein, the term "staple length fibers" refers to those fibers having a length of at least about 15.9 mm (0.625 inches). Preferably, the topsheet has a basis weight from about 14 to about 25 grams per square meter. A suitable topsheet is manufactured by Veratec, Inc., a Division of International Paper 30 Company, of Walpole, Mass. under the designation P-8.

The topsheet 520 of diaper 50 is preferably made of a hydrophilic material to promote rapid transfer of liquids (e.g., urine) through the topsheet. If the topsheet is made of a hydrophobic material, at least the upper surface of the topsheet is treated to be hydrophilic so that liquids will transfer through the topsheet more rapidly. This diminishes the likelihood 35 that body fluids will flow off the topsheet rather than being drawn

through the topsheet and being absorbed by the absorbent core. The topsheet can be rendered hydrophilic by treating it with a surfactant. Suitable methods for treating the topsheet with a surfactant include spraying the topsheet material with the surfactant and immersing the material into the surfactant. A more detailed discussion of such a treatment and hydrophilicity is contained in U.S. Patents 4,988,344 entitled "Absorbent Articles with Multiple Layer Absorbent Layers" issued to Reising, et al on January 29, 1991 and U.S. Patent 4,988,345 entitled "Absorbent Articles with Rapid Acquiring Absorbent Cores" issued to Reising on January 29, 1991, each of which is incorporated by reference herein.

5 In a preferred embodiment of a diaper as described herein, the backsheet 530 has a modified hourglass shape extending beyond the absorbent core a minimum distance of about 1.3 cm to about 6.4 cm (about 10 0.5 to about 2.5 inch) around the entire diaper periphery.

15 The absorbent core 540 may take on any size or shape that is compatible with the diaper 50. One preferred embodiment of the diaper 50 has an asymmetric, modified T-shaped absorbent core 540 having ears in the first waist region but a generally rectangular shape in the second waist region. Exemplary absorbent structures for use as the absorbent core of the present invention that have achieved wide acceptance and commercial success are described in U.S. Patent 4,610,678 entitled "High-Density Absorbent Structures" issued to Weisman et al. on September 9, 1986; U.S. Patent 4,673,402 entitled "Absorbent Articles With Dual-Layered Cores" 20 issued to Weisman et al. on June 16, 1987; U.S. Patent 4,888,231 entitled "Absorbent Core Having A Dusting Layer" issued to Angstadt on December 19, 1989; and U.S. Patent 4,834,735, entitled "High Density Absorbent Members Having Lower Density and Lower Basis Weight Acquisition Zones", issued to Alemany et al. on May 30, 1989. The absorbent core 25 may further comprise the dual core system containing an acquisition/distribution core of chemically stiffened fibers positioned over an absorbent storage core as detailed in U.S. Patent 5,234,423, entitled "Absorbent Article With Elastic Waist Feature and Enhanced Absorbency" issued to Alemany et al., on August 10, 1993; and in U.S. Patent 5,147,345, 30 entitled "High Efficiency Absorbent Articles For Incontinence Management" 35

issued to Young, LaVon and Taylor on September 15, 1992. All of these patents are incorporated herein by reference.

In a preferred embodiment, the diaper 50 further comprises elasticized leg cuffs 550 for providing improved containment of liquids and other body exudates; an elastic waist feature 560 that provides improved fit and containment; and a fastening system 570 which forms a side closure which maintains the first waist region 56 and the second waist region 58 in an overlapping configuration such that lateral tensions are maintained around the circumference of the diaper to maintain the diaper on the wearer.

5 The diaper 50 may also comprise elasticized side panels (not shown) in the waist regions 56 and 58 to provide an elastically extensible feature that provides a more comfortable and contouring fit and more effective application of the diaper 50.

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The elasticized leg cuffs 550 can be constructed in a number of different configurations, including those described in U.S. Patent No. 3,860,003; U.S. Patent No. 4,909,803, issued to Aziz et al. on Mar. 20, 1990; U.S. Patent No. 4,695,278, issued to Lawson on Sep. 22, 1987; and U.S. Patent No. 4,795,454, issued to Dragoo on Jan. 3, 1989, each being incorporated herein by reference.

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20 The elasticized waist feature preferably comprises an elasticized waistband (not shown) that may be constructed in a number of different configurations including those described in U.S. Patent No. 4,515,595 issued to Kievit et al. on May 7, 1985; U.S. Patent No. 5,026,364 issued to Robertson on Jun. 25, 1991; and the above referenced U.S. Patent No. 25 5,151,092 issued to Buell et al. on Sep. 29, 1992, each of these references being incorporated herein by reference.

20 The elasticized side panels may be constructed in a number of configurations. Examples of diapers with elasticized side panels positioned in the ears (ear flaps) of the diaper are disclosed in U.S. Patent No. 30 4,857,067, issued to Wood, et al. on Aug. 15, 1989; U.S. Patent No. 4,381,781, issued to Sciarappa, et al. on May 3, 1983; U.S. Patent No. 4,938,753, issued to Van Gompel, et al. on Jul. 3, 1990; and U.S. Patent No. 5,151,092, issued to Buell et al. on Sep. 29, 1992; each of which are incorporated herein by reference.

Exemplary fastening systems 570 are disclosed in U.S. Patent No. 4,846,815, issued to Scripps on Jul. 11, 1989; U.S. Patent No. 4,894,060, issued to Nestegard on Jan. 16, 1990; U.S. Patent No. 4,946,527, issued to Battrell on Aug. 7, 1990; U.S. Patent No. 3,848,594, issued to Buell on Nov. 5, 1974; U.S. Patent No. B1 4,662,875, issued to Hirotsu et al. on May 5, 1987; and U.S. Patent No. 5,151,092, issued to Buell et al. on Sep. 29, 1992; each of which is incorporated herein by reference.

The diaper 50 is preferably applied to a wearer by positioning one of the waist regions of the diaper, preferably the second waist region 58, under the wearer's back and drawing the remainder of the diaper between the wearer's legs so that the other waist region, preferably the first waist region 56, is positioned across the front of the wearer. The fastening system is then applied to effect a side closure.

The lotioned topsheets of the present invention are also useful in training pants. The term "training pants", as used herein, refers to disposable garments having fixed sides and leg openings. Training pants are placed in position on the wearer by inserting the wearer's legs into the leg openings and sliding the training pant into position about the wearer's lower torso. Suitable training pants are disclosed in U.S. Patent No. 5,246,433, issued to Hasse, et al. on September 21, 1993.

Another disposable absorbent article for which the lotioned topsheets of the present invention are useful are incontinence articles. The term "incontinence article" refers to pads, undergarments (pads held in place by a suspension system of same type, such as a belt, or the like), inserts for absorbent articles, capacity boosters for absorbent articles, briefs, bed pads, and the like regardless of whether they are worn by adults or other incontinent persons. Suitable incontinence articles are disclosed in U.S. Patent No. 4,253,461 issued to Strickland, et al. on March 3, 1981; U.S. Patent Nos. 4,597,760 and 4,597,761 issued to Buell; the above-mentioned U.S. Patent No. 4,704,115; U.S. Patent No. 4,909,802 issued to Ahr, et al.; U.S. Patent No. 4,964,860 issued to Gipson, et al. on October 23, 1990; and in U.S. Patent Application Serial No. 07/637,090 filed by Noel, et al. on January 3, 1991 (PCT Publication No. WO 92/11830 published on July 23, 1992).

B. Lotion Composition.

The lotion compositions of the present invention are solid, or more often semisolid, at 20°C, i.e. at ambient temperatures. By "semisolid" is meant that the lotion composition has a rheology typical of pseudoplastic or plastic fluids. When no shear is applied, the lotion compositions can have the appearance of a semi-solid but can be made to flow as the shear rate is increased. This is due to the fact that, while the lotion composition contains primarily solid components, it also includes some minor liquid components.

The lotion compositions of the present invention are at least semi-solid at room temperature to minimize lotion migration. In addition, the lotion compositions preferably have a final melting point (100% liquid) above potential "stressful" storage conditions that can be greater than 45°C (e.g., warehouse in Arizona, car trunk in Florida, etc.).

Specifically, the lotion compositions of the present invention should have the following melt profile:

<u>Characteristic</u>	<u>Preferred Range</u>	<u>Most Preferred</u>
% liquid at room temp. (20 °C)	2-50	3-25
% liquid at body temp. (37 °C)	25-95	30-90
final melting point (°C)	≥38	≥45

By being solid or semisolid at ambient temperatures, these lotion compositions do not have a tendency to flow and migrate into the interior of the diaper topsheet to which they are applied. This means less lotion composition is required for imparting desirable therapeutic or protective coating lotion benefits.

When applied to outer surface of diaper topsheets, the lotion compositions of the present invention are transferable to the wearer's skin by normal contact, wearer motion, and/or body heat. Importantly, the lotions disclosed in the present invention reduce the adherence of BM to the skin of the wearer, thereby improving the ease of BM clean up.

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CF	Central African Republic	KP	Democratic People's Republic of Korea	SD	Sudan
CG	Congo	KR	Republic of Korea	SE	Sweden
CH	Switzerland	KZ	Kazakhstan	SI	Slovenia
CI	Côte d'Ivoire	LI	Liechtenstein	SK	Slovakia
CM	Cameroon	LK	Sri Lanka	SN	Senegal
CN	China	LU	Luxembourg	TD	Chad
CS	Czechoslovakia	LV	Latvia	TG	Togo
CZ	Czech Republic	MC	Monaco	TJ	Tajikistan
DE	Germany	MD	Republic of Moldova	TT	Trinidad and Tobago
DK	Denmark	MG	Madagascar	UA	Ukraine
ES	Spain	ML	Mali	US	United States of America
FI	Finland	MN	Mongolia	UZ	Uzbekistan
FR	France			VN	Viet Nam
GA	Gabon				

for example, ambient humidity can be tolerated without adverse effect. Typically, the emollients used in the present invention contain about 5% or less water, preferably about 1% or less water, most preferably about 0.5% or less water.

5 Emollients useful in the present invention can be petroleum-based, fatty acid ester type, alkyl ethoxylate type, fatty acid ester ethoxylates, fatty alcohol type, polysiloxane type, or mixtures of these emollients. Suitable petroleum-based emollients include those hydrocarbons, or mixtures of hydrocarbons, having chain lengths of from 16 to 32 carbon atoms.

10 Petroleum based hydrocarbons having these chain lengths include mineral oil (also known as "liquid petrolatum") and petrolatum (also known as "mineral wax," "petroleum jelly" and "mineral jelly"). Mineral oil usually refers to less viscous mixtures of hydrocarbons having from 16 to 20 carbon atoms. Petrolatum usually refers to more viscous mixtures of hydrocarbons

15 having from 16 to 32 carbon atoms. Petrolatum and mineral oil are particularly preferred emollients for lotion compositions of the present invention.

20 Suitable fatty acid ester type emollients include those derived from C₁₂-C₂₈ fatty acids, preferably C₁₆-C₂₂ saturated fatty acids, and short chain (C₁-C₈, preferably C₁-C₃) monohydric alcohols. Representative examples of such esters include methyl palmitate, methyl stearate, isopropyl laurate, isopropyl myristate, isopropyl palmitate, ethylhexyl palmitate and mixtures thereof. Suitable fatty acid ester emollients can also be derived from esters of longer chain fatty alcohols (C₁₂-C₂₈, preferably C₁₂-C₁₆) and shorter chain fatty acids e.g., lactic acid, such as lauryl lactate and cetyl lactate.

25 Suitable alkyl ethoxylate type emollients include C₁₂-C₂₂ fatty alcohol ethoxylates having an average degree of ethoxylation of from about 2 to about 30. Preferably, the fatty alcohol ethoxylate emollient is selected from the group consisting of lauryl, cetyl, and stearyl ethoxylates, and mixtures thereof, having an average degree of ethoxylation ranging from about 2 to about 23. Representative examples of such alkyl ethoxylates include laureth-3 (a lauryl ethoxylate having an average degree of ethoxylation of 3), laureth-23 (a lauryl ethoxylate having an average degree of ethoxylation of 23), cet th-10 (a cetyl alcohol ethoxylate having an average degree of ethoxylation of 10) and st areth-10 (a stearyl alcohol

ethoxylate having an average degree of ethoxylation of 10). These alkyl ethoxylate emollients are typically used in combination with the petroleum-based emollients, such as petrolatum, at a weight ratio of alkyl ethoxylate emollient to petroleum-based emollient of from about 1:1 to about 1:5, 5 preferably from about 1:2 to about 1:4.

Suitable fatty alcohol type emollients include C₁₂-C₂₂ fatty alcohols, 10 preferably C₁₆-C₁₈ fatty alcohols. Representative examples include cetyl alcohol and stearyl alcohol, and mixtures thereof. These fatty alcohol emollients are typically used in combination with the petroleum-based emollients, such as petrolatum, at a weight ratio of fatty alcohol emollient to petroleum-based emollient of from about 1:1 to about 1:5, preferably from 15 about 1:1 to about 1:2.

Other suitable types of emollients for use in the present invention include polysiloxane compounds. In general suitable polysiloxane materials for use in the present invention include those having monomeric siloxane units of the following structure:

R₁

(1) - Si - O-

R₂

wherein, R₁ and R₂, for each independent siloxane monomeric unit can 20 each independently be hydrogen or any alkyl, aryl, alkenyl, alkaryl, arakyl, cycloalkyl, halogenated hydrocarbon, or other radical. Any of such radicals can be substituted or unsubstituted. R₁ and R₂ radicals of any particular monomeric unit may differ from the corresponding functionalities of the next adjoining monomeric unit. Additionally, the polysiloxane can be either a straight chain, a branched chain or have a cyclic structure. The radicals R₁ and R₂ can additionally independently be other silaceous functionalities 25 such as, but not limited to siloxanes, polysiloxanes, silanes, and polysilanes. The radicals R₁ and R₂ may contain any of a variety of 30 organic functionaliti s including, for exempl , alcohol, carboxylic acid, ph nyt, and amine functionaliti s.

Exemplary alkyl radicals are methyl, ethyl, propyl, butyl, pentyl, hexyl; octyl, decyl, octadecyl, and the like. Exemplary alkenyl radicals are vinyl, allyl, and the like. Exemplary aryl radicals are phenyl, diphenyl, naphthyl, and the like. Exemplary alkaryl radicals are toyl, xylyl, ethylphenyl, and the like. Exemplary aralkyl radicals are benzyl, alpha-phenylethyl, beta-phenylethyl, alpha-phenylbutyl, and the like. Exemplary cycloalkyl radicals are cyclobutyl, cyclopentyl, cyclohexyl, and the like. Exemplary halogenated hydrocarbon radicals are chloromethyl, bromoethyl, tetrafluoroethyl, fluoreethyl, trifluoreethyl, trifluorotioyl, hexafluoroxylyl, and the like.

Viscosity of polysiloxanes useful may vary as widely as the viscosity of polysiloxanes in general vary, so long as the polysiloxane is flowable or can be made to be flowable for application to the diaper topsheet. This includes, but is not limited to, viscosity as low as 5 centistokes (at 37°C as measured by a glass viscometer) to about 20,000,000 centistokes. Preferably the polysiloxanes have a viscosity at 37°C ranging from about 5 to about 5,000 centistokes, more preferably from about 5 to about 2,000 centistokes, most preferably from about 100 to about 1000 centistokes. High viscosity polysiloxanes which themselves are resistant to flowing can be effectively deposited upon the diaper topsheets by such methods as, for example, emulsifying the polysiloxane in surfactant or providing the polysiloxane in solution with the aid of a solvent, such as hexane, listed for exemplary purposes only. Particular methods for applying polysiloxane emollients to diaper topsheets are discussed in more detail hereinafter.

Preferred polysiloxanes compounds for use in the present invention are disclosed in U.S. Patent 5,059,282 (Ampulski et al), issued October 22, 1991, which is incorporated herein by reference. Particularly preferred polysiloxane compounds for use as emollients in the lotion compositions of the present invention include phenyl-functional polymethylsiloxane compounds (e.g., Dow Corning 556 Cosmetic-Grade Fluid: polyphenylmethylsiloxane) and cetyl or stearyl functionalized dimethicones such as Dow 2502 and Dow 2503 polysiloxane fluids, respectively. In addition to such substitution with phenyl-functional or alkyl groups, effective substitution may be made with amino, carboxyl, hydroxyl, ether, poly thio, aldehyd, ketone, amid, ester, and thiol groups. Of these effective substituent groups, the family of groups comprising phenyl, amino,

alkyl, carboxyl, and hydroxyl groups are more preferred than the others; and phenyl-functional groups are most preferred.

5 Besides petroleum-based emollients, fatty acid ester emollients, fatty acid ester ethoxylates, alkyl ethoxylate emollients fatty alcohol emollients, and polysiloxanes, the emollients useful in the present invention can include minor amounts (e.g., up to about 10% of the total emollient) of other, conventional emollients. These other, conventional emollients include propylene glycol, glycerine, triethylene glycol, spermaceti or other waxes, fatty acids; and fatty alcohol ethers having from 12 to 28 carbon atoms in their fatty chain, such as stearic acid, propoxylated fatty alcohols; glycerides, acetoglycerides, and ethoxylated glycerides of C₁₂-C₂₈ fatty acids; other fatty esters of polyhydroxy alcohols; lanolin and its derivatives. These other emollients should be included in a manner such that the solid or semisolid characteristics of the lotion composition are maintained.

10 15 The amount of emollient that can be included in the lotion composition will depend on a variety of factors, including the particular emollient involved, the lotion-like benefits desired, the other components in the lotion composition and like factors. The lotion composition can comprise from about 10 to about 95% of the emollient. Preferably, the lotion composition comprises from about 20 to about 80%, most preferably from about 40 to about 75%, of the emollient.

20 25 30 35

2. Immobilizing Agent

An especially key component of the lotion compositions of the present invention is an agent capable of immobilizing the emollient on the diaper topsheet to which the lotion composition is applied. Because the emollient in the composition has a plastic or fluid consistency at 20°C, it tends to flow or migrate, even when subjected to modest shear. When applied to a diaper topsheet, especially in a melted or molten state, the emollient will not remain primarily on the surface of the topsheet. Instead, the emollient will tend to migrate and flow into the interior of the diaper.

This migration of the emollient into the interior of the diaper can cause undesired effects on the absorbency of the diaper core due to the hydrophobic characteristics of many of the emollients used in the lotion compositions of the present invention. It also means that much more

emollient has to be applied to the diaper topsheet to get the desired therapeutic or protective lotion benefits. Increasing the level of emollient not only increases the cost, but also exacerbates the undesirable effect on the absorbency of the diaper core.

5 The immobilizing agent counteracts this tendency of the emollient to migrate or flow by keeping the emollient primarily localized on the surface of the diaper topsheet to which the lotion composition is applied. This is believed to be due, in part, to the fact that the immobilizing agent raises the melting point of the lotion composition above that of the emollient. Since
10 the immobilizing agent is also miscible with the emollient (or solubilized in the emollient with the aid of an appropriate emulsifier), it entraps the emollient on the surface of the diaper topsheet as well.

15 It is also advantageous to "lock" the immobilizing agent on the surface of the diaper topsheet. This can be accomplished by using immobilizing agents which quickly crystallize (i.e., solidify) at the surface of the topsheet. In addition, outside cooling of the treated diaper topsheet via blowers, fans, etc. can speed up crystallization of the immobilizing agent.

20 In addition to being miscible with (or solubilized in) the emollient, the immobilizing agent needs to have a melting point of at least about 35°C. This is so the immobilizing agent itself will not have a tendency to migrate or flow. Preferred immobilizing agents will have melting points of at least about 40°C. Typically, the immobilizing agent will have a melting point in the range of from about 50° to about 150°C.

25 Suitable immobilizing agents for the present invention can comprise a member selected from the group consisting of C₁₄-C₂₂ fatty alcohols, C₁₂-C₂₂ fatty acids, and C₁₂-C₂₂ fatty alcohol ethoxylates having an average degree of ethoxylation ranging from 2 to about 30, and mixtures thereof. Preferred immobilizing agents include C₁₆-C₁₈ fatty alcohols, most preferably selected from the group consisting of cetyl alcohol, stearyl alcohol, and mixtures thereof. Mixtures of cetyl alcohol and stearyl alcohol are particularly preferred. Other preferred immobilizing agents include C₁₆-C₁₈ fatty acids, most preferably selected from the group consisting of palmitic acid, stearic acid, and mixtures thereof. Mixtures of palmitic acid and stearic acid are particularly preferred. Still other preferred immobilizing agents include C₁₆-C₁₈ fatty alcohol ethoxylates having an average

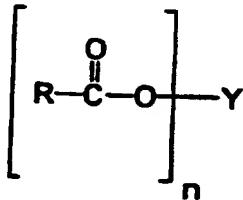
degree of ethoxylation ranging from about 5 to about 20. Preferably, the fatty alcohols, fatty acids and fatty alcohols are linear.

Importantly, these preferred immobilizing agents such as the C₁₆ - C₁₈ fatty alcohols increase the rate of crystallization of the lotion causing the lotion to crystallize rapidly onto the surface of the substrate. Lower lotion levels can therefore be utilized or a superior lotion feel can be delivered. Traditionally, greater amounts of lotion were needed to generate softness because of the flow of these liquids into the diaper core.

Other types of immobilizing agents can be used either alone or in combination with the fatty alcohols, fatty acids, and fatty alcohol ethoxylates described above. Examples of these other types of immobilizing agents includes polyhydroxy fatty acid esters, polyhydroxy fatty acid amides, and mixtures thereof. Preferred esters and amides will have three or more free hydroxy groups on the polyhydroxy moiety and are typically nonionic in character. Because of the possible skin sensitivity of those using diaper topsheets to which the lotion composition is applied, these esters and amides should also be relatively mild and non-irritating to the skin.

Suitable polyhydroxy fatty acid esters for use in the present invention will have the formula:

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wherein R is a C₅-C₃₁ hydrocarbyl group, preferably straight chain C₇-C₁₉ alkyl or alkenyl, more preferably straight chain C₉-C₁₇ alkyl or alkenyl, most preferably straight chain C₁₁-C₁₇ alkyl or alkenyl, or mixture thereof; Y is a polyhydroxyhydrocarbyl moiety having a hydrocarbyl chain with at least 2 free hydroxyls directly connected to the chain; and n is at least 1. Suitable Y groups can be derived from polyols such as glycerol, pentaerythritol; sugars such as raffinose, maltodextrose, galactose, sucrose, glucose, xylose, fructose, maltose, lactose, mannose and

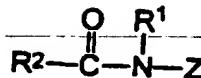
erythrose; sugar alcohols such as erythritol, xylitol, malitol, mannitol and sorbitol; and anhydrides of sugar alcohols such as sorbitan.

One class of suitable polyhydroxy fatty acid esters for use in the present invention comprises certain sorbitan esters, preferably the sorbitan esters of C₁₆-C₂₂ saturated fatty acids. Because of the manner in which they are typically manufactured, these sorbitan esters usually comprise mixtures of mono-, di-, tri-, etc. esters. Representative examples of suitable sorbitan esters include sorbitan palmitates (e.g., SPAN 40), sorbitan stearates (e.g., SPAN 60), and sorbitan behenates, that comprise one or 5 more of the mono-, di- and tri-ester versions of these sorbitan esters, e.g., sorbitan mono-, di- and tri-palmitate, sorbitan mono-, di- and tri-stearate, sorbitan mono-, di- and tri-behenate, as well as mixed tallow fatty acid sorbitan mono-, di- and tri-esters. Mixtures of different sorbitan esters can 10 also be used, such as sorbitan palmitates with sorbitan stearates. 15 Particularly preferred sorbitan esters are the sorbitan stearates, typically as a mixture of mono-, di- and tri-esters (plus some tetraester) such as SPAN 60, and sorbitan stearates sold under the trade name GLYCOMUL-S by Lonza, Inc. Although these sorbitan esters typically contain mixtures of 20 mono-, di- and tri-esters, plus some tetraester, the mono- and di-esters are usually the predominant species in these mixtures.

Another class of suitable polyhydroxy fatty acid esters for use in the present invention comprises certain glyceryl monoesters, preferably glyceryl monoesters of C₁₆-C₂₂ saturated fatty acids such as glyceryl monostearate, glyceryl monopalmitate, and glyceryl monobehenate. Again, 25 like the sorbitan esters, glyceryl monoester mixtures will typically contain some di- and triester. However, such mixtures should contain predominantly the glyceryl monoester species to be useful in the present invention.

Another class of suitable polyhydroxy fatty acid esters for use in the 30 present invention comprise certain sucrose fatty acid esters, preferably the C₁₂-C₂₂ saturated fatty acid esters of sucrose. Sucrose monoesters and diesters are particularly preferred and include sucrose mono- and di-stearate and sucrose mono- and di-laureate.

Suitable polyhydroxy fatty acid amids for use in the present 35 invention will have the formula:



5 wherein R^1 is H, C₁-C₄ hydrocarbyl, 2-hydroxyethyl, 2-hydroxypropyl, methoxyethyl, methoxypropyl or a mixture thereof, preferably C₁-C₄ alkyl, methoxyethyl or methoxypropyl, more preferably C₁ or C₂ alkyl or methoxypropyl, most preferably C₁ alkyl (i.e., methyl) or methoxypropyl; and R^2 is a C₅-C₃₁ hydrocarbyl group, preferably straight chain C₇-C₁₉ alkyl or alkenyl, more preferably straight chain C₉-C₁₇ alkyl or alkenyl, most preferably straight chain C₁₁-C₁₇ alkyl or alkenyl, or mixture thereof; 10 and Z is a polyhydroxyhydrocarbyl moiety having a linear hydrocarbyl chain with at least 3 hydroxyls directly connected to the chain. See U.S. patent 5,174, 927 (Honsa), issued December 29, 1992 (herein incorporated by reference) which discloses these polyhydroxy fatty acid amides, as well as 15 their preparation.

The Z moiety preferably will be derived from a reducing sugar in a reductive amination reaction; most preferably glycetyl. Suitable reducing sugars include glucose, fructose, maltose, lactose, galactose, mannose, and xylose. High dextrose corn syrup, high fructose corn syrup, and high 20 maltose corn syrup can be utilized, as well as the individual sugars listed above. These corn syrups can yield mixtures of sugar components for the Z moiety.

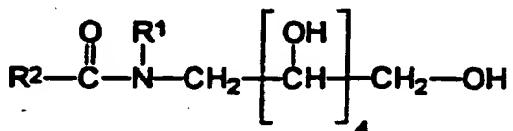
The Z moiety preferably will be selected from the group consisting of -CH₂-(CHOH)_n-CH₂OH, -CH(CH₂OH)-[(CHOH)_{n-1}]-CH₂OH, -CH₂OH-CH₂-(CHOH)₂(CHOR³)(CHOH)-CH₂OH, where n is an integer from 3 to 5, 25 and R³ is H or a cyclic or aliphatic monosaccharide. Most preferred are the glycetyl where n is 4, particularly -CH₂-(CHOH)₄-CH₂OH.

In the above formula, R¹ can be, for example, N-methyl, N-ethyl, N-propyl, N-isopropyl, N-butyl, N-2-hydroxyethyl, N-methoxypropyl or N-2-hydroxypropyl. R² can be selected to provide, for example, cocamides, 30 stearamides, oleamides, lauramides, myristamides, capricamides, palmitamides, tallowamides, etc. The Z moiety can be 1-deoxyglucetyl, 2-

deoxyfructityl, 1-deoxymaltityl, 1-deoxylactityl, 1-deoxygalactityl, 1-deoxymannityl, 1-deoxymaltotriotityl, etc.

The most preferred polyhydroxy fatty acid amides have the general formula:

5



wherein R^1 is methyl or methoxypropyl; R^2 is a C_{11} - C_{17} straight-chain alkyl or alkenyl group. These include N-lauryl-N-methyl glucamide, N-lauryl-N-methoxypropyl glucamide, N-cocoyl-N-methyl glucamide, N-cocoyl-N-methoxypropyl glucamide, N-palmityl-N-methoxypropyl glucamide, N-tallowyl-N-methyl glucamide, or N-tallowyl-N-methoxypropyl glucamide.

As previously noted, some of the immobilizing agents require an emulsifier for solubilization in the emollient. This is particularly the case for certain of the glucamides such as the N-alkyl-N-methoxypropyl glucamides having HLB values of at least about 7. Suitable emulsifiers will typically include those having HLB values below about 7. In this regard, the sorbitan esters previously described, such as the sorbitan stearates, having HLB values of about 4.9 or less have been found useful in solubilizing these glucamide immobilizing agents in petrolatum. Other suitable emulsifiers include steareth-2 (polyethylene glycol ethers of stearyl alcohol that conform to the formula $\text{CH}_3(\text{CH}_2)_{17}(\text{OCH}_2\text{CH}_2)_n\text{OH}$, where n has an average value of 2), sorbitan tristearate, isosorbide laurate, and glyceryl monostearate. The emulsifier can be included in an amount sufficient to solubilize the immobilizing agent in the emollient such that a substantially homogeneous mixture is obtained. For example, an approximately 1:1 mixture of N-cocoyl-N-methyl glucamide and petrolatum that will normally not melt into a single phase mixture, will melt into a single phase mixture upon the addition of 20% of a 1:1 mixture of Steareth-2 and sorbitan tristearate as the emulsifier.

Other types of ingredients that can be used as immobilizing agents, either alone, or in combination with the above-mentioned immobilizing

agents, include waxes such as carnauba, beeswax, candelilla, paraffin, ceresin, esparto, ouricuri, rezowax, and other known waxes. Preferably the wax is a paraffin wax. An example of a particularly preferred paraffin wax is Paraffin S.P. 434 from Strahl and Pitsch Inc. P.O. Box 1098 West Babylon, 5 NY 11704.

10 The amount of immobilizing agent that should be included in the lotion composition will depend on a variety of factors, including the particular emollient involved, the particular immobilizing agent involved, whether an emulsifier is required to solubilize the immobilizing agent in the emollient, the other components in the lotion composition and like factors. The lotion composition can comprise from about 5 to about 90% of the immobilizing agent. Preferably, the lotion composition comprises from about 5 to about 50%, most preferably from about 10 to about 40%, of the immobilizing agent.

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3. Optional Hydrophilic Surfactant

20 As mentioned above, it is highly desirable that the diaper topsheet is made of a hydrophilic material to promote rapid transfer of liquids (e.g., urine) through the topsheet. Similarly, it is important that the lotion composition also be sufficiently wettable to ensure that liquids will transfer through the topsheet more rapidly. This diminishes the likelihood that body exudates will flow off the lotion coating rather than being drawn through the topsheet and being absorbed by the absorbent core. Depending upon the 25 particular immobilizing agent used in the lotion composition of the present invention, an additional hydrophilic surfactant (or a mixture of hydrophilic surfactants) may, or may not, be required to improve wettability. For example, some immobilizing agents, such as N-cocoyl-N-methoxypropyl glucamide have HLB values of at least about 7 and are sufficiently wettable without the addition of hydrophilic surfactant. Other immobilizing agents 30 such as the C₁₆ - C₁₈ fatty alcohols having HLB values below about 7 will require addition of hydrophilic surfactant to improve wettability when the lotion composition is applied to diaper topsheets. Similarly, a hydrophobic emollient such as petrolatum will require the addition of a hydrophilic surfactant.

Suitable hydrophilic surfactants will be miscible with the emollient and the immobilizing agent so as to form homogeneous mixtures. Because of possible skin sensitivity of those using disposable absorbent products to which the lotion composition is applied, these surfactants should also be 5 relatively mild and non-irritating to the skin. Typically, these hydrophilic surfactants are nonionic to be not only non-irritating to the skin, but also to avoid other undesirable effects on any underlying tissue laminate structure, e.g., reductions in tensile strength.

Suitable nonionic surfactants may be substantially nonmigratory after 10 the lotion composition is applied to the diaper topsheets and will typically have HLB values in the range of from about 4 to about 20, preferably from about 7 to about 20. To be nonmigratory, these nonionic surfactants will typically have melt temperatures greater than the temperatures commonly encountered during storage, shipping, merchandising, and use of 15 disposable absorbent products, e.g., at least about 30°C In this regard, these nonionic surfactants will preferably have melting points similar to those of the immobilizing agents previously described.

Suitable nonionic surfactants for use in lotion compositions of the 20 present invention include alkylglycosides; alkylglycoside ethers as described in U.S. patent 4,011,389 (Langdon, et al), issued March 8, 1977; alkylpolyethoxylated esters such as Pegosperse 1000MS (available from Lonza, Inc., Fair Lawn, New Jersey), ethoxylated sorbitan mono-, di- and/or tri-esters of C₁₂-C₁₈ fatty acids having an average degree of ethoxylation of from about 2 to about 20, preferably from about 2 to about 10, such as 25 TWEEN 60 (sorbitan esters of stearic acid having an average degree of ethoxylation of about 20) and TWEEN 61 (sorbitan esters of stearic acid having an average degree of ethoxylation of about 4), and the condensation products of aliphatic alcohols with from about 1 to about 54 moles of ethylene oxide. The alkyl chain of the aliphatic alcohol is typically in a 30 straight chain (linear) configuration and contains from about 8 to about 22 carbon atoms. Particularly preferred are the condensation products of alcohols having an alkyl group containing from about 11 to about 22 carbon atoms with from about 2 to about 30 moles of ethylene oxide per mole of alcohol. Examples of such ethoxylated alcohols include the condensation 35 products of myristyl alcohol with 7 moles of thylene oxid per m l of alcohol, the condensation products of coconut alcohol (a mixtur of fatty

alcohols having alkyl chains varying in length from 10 to 14 carbon atoms) with about 6 moles of ethylene oxide. A number of suitable ethoxylated alcohols are commercially available, including TERGITOL 15-S-9 (the condensation product of C₁₁-C₁₅ linear alcohols with 9 moles of ethylene oxide), marketed by Union Carbide Corporation; KYRO EOB (condensation product of C₁₃-C₁₅ linear alcohols with 9 moles of ethylene oxide), marketed by The Procter & Gamble Co., the NEODOL brand name surfactants marketed by Shell Chemical Co., in particular NEODOL 25-12 (condensation product of C₁₂-C₁₅ linear alcohols with 12 moles of ethylene oxide) and NEODOL 23-6.5T (condensation product of C₁₂-C₁₃ linear alcohols with 6.5 moles of ethylene oxide that has been distilled (topped) to remove certain impurities), and especially the PLURAFAC brand name surfactants marketed by BASF Corp., in particular PLURAFAC A-38 (a condensation product of a C₁₈ straight chain alcohol with 27 moles of ethylene oxide). (Certain of the hydrophilic surfactants, in particular ethoxylated alcohols such as NEODOL 25-12, can also function as alkyl ethoxylate emollients). Other examples of preferred ethoxylated alcohol surfactants include ICI's class of Brij surfactants and mixtures thereof, with Brij 72 (i.e., Steareth-2) and Brij 76 (i.e., Steareth-10) being especially preferred. Also, mixtures of cetyl alcohol and stearyl alcohol ethoxylated to an average degree of ethoxylation of from about 10 to about 20 may also be used as the hydrophilic surfactant.

Another type of suitable surfactant for use in the present invention includes Aerosol OT, a dioctyl ester of sodium sulfosuccinic acid marketed by American Cyanamid Company.

Still another type of suitable surfactant for use in the present invention includes silicone copolymers such as General Electric SF 1188 (a copolymer of a polydimethylsiloxane and a polyoxyalkylene ether) and General Electric SF 1228 (a silicone polyether copolymer). These silicone surfactants can be used in combination with the other types of hydrophilic surfactants discussed above, such as the ethoxylated alcohols. These silicone surfactants have been found to be effective at concentrations as low as 0.1%, more preferably from about 0.25 to about 1.0%, by weight of the lotion composition.

The amount of hydrophilic surfactant required to increase the wettability of the lotion composition to a desired level will depend upon the

HLB value and level of immobilizing agent used, the HLB value of the surfactant used and like factors. The lotion composition can comprise from about 1 to about 50% of the hydrophilic surfactant when needed to increase the wettability properties of the composition. Preferably, the lotion 5 composition comprises from about 1 to about 25%, most preferably from about 10 to about 20%, of the hydrophilic surfactant when needed to increase wettability.

4. Other Optional Components

Lotion compositions can comprise other optional components 10 typically present in emollient, creams, and lotions of this type. These optional components include water, viscosity modifiers, perfumes, disinfectant antibacterial actives, pharmaceutical actives, film formers, deodorants, opacifiers, astringents, solvents and the like. In addition, stabilizers can be added to enhance the shelf life of the lotion composition 15 such as cellulose derivatives, proteins and lecithin. All of these materials are well known in the art as additives for such formulations and can be employed in appropriate amounts in the lotion compositions of the present invention.

20 C. Treating Diaper Topsheets With Lotion Composition

In preparing lotioned diaper products according to the present invention, the lotion composition is applied to the outer surface (i.e., body facing surface) of a diaper topsheet. Any of a variety of application methods that evenly distribute lubricious materials having a molten or liquid 25 consistency can be used. Suitable methods include spraying, printing (e.g., flexographic printing), coating (e.g., gravure coating), extrusion, or combinations of these application techniques, e.g. spraying the lotion composition on a rotating surface, such as a calender roll, that then transfers the composition to the outer surface of the diaper topsheet.

30 The manner of applying the lotion composition to the diaper topsheet should be such that the topsheet does not become saturated with the lotion composition. If the topsheet becomes saturated with the lotion composition, there is a greater potential for the lotion to block the topsheet openings, reducing the ability of the topsheet to transmit fluid to the underlying 35 absorbent core. Also, saturation of the topsheet is not required to obtain

the therapeutic and/or protective lotion benefits. Particularly suitable application methods will apply the lotion composition primarily to the outer surface of the diaper topsheet.

5 The minimum level of lotion to be applied to the diaper topsheets is an amount effective for reducing the adherence of BM to the skin of the wearer. The lotion composition is preferably applied to the diaper topsheets of the present invention in an amount ranging from about 0.1 mg/in² to about 25 mg/in² more preferably from about 1 mg/in² to about 10 mg/in² (mg of lotion per square inch of coated topsheet). Because the emollient is substantially immobilized on the surface of the topsheet, less lotion composition is needed to impart the desired skin care benefits. Such relatively low levels of lotion composition are adequate to impart the desired therapeutic and/or protective lotion benefits to the topsheet, yet do not saturate the topsheet's absorbency and/or wettability properties.

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The lotion composition may be applied to the entire surface of the topsheet or portions thereof. Preferably, the lotion composition is applied in a stripe aligned with and centered on the longitudinal centerline of the disposable absorbent article. Most preferably, as described in the Examples hereinafter, the lotion composition is applied to a discrete version of the topsheet, e.g., a 3.75 inch wide (diaper lateral direction) and 7 inch long (diaper longitudinal direction) patch generally disposed toward the second waist region.

The lotion composition can also be applied nonuniformly to the outer surface of the diaper topsheet. By "nonuniform" is meant that the amount, pattern of distribution, etc. of the lotion composition can vary over the topsheet surface. For example, some portions of the treated surface of the topsheet can have greater or lesser amounts of lotion composition, including portions of the surface that do not have any lotion composition on it.

The lotion composition can be applied to the topsheet at any point during assembly. For example, the lotion composition can be applied to the topsheet of the finished disposable absorbent product before it has been packaged. The lotion composition can also be applied to the topsheet before it is combined with the other raw materials to form a finished disposable absorbent product.

The lotion composition is typically applied from a melt thereof to the diaper topsheet. Since the lotion composition melts at significantly above ambient temperatures, it is usually applied as a heated coating to-the topsheet. Typically, the lotion composition is heated to a temperature in the 5 range from about 35° to about 100°C, preferably from 40° to about 90°C, prior to being applied to the diaper topsheet. Once the melted lotion composition has been applied to the diaper topsheet, it is allowed to cool and solidify to form solidified coating or film on the surface of the topsheet. Preferably, the application process is designed to aid in the cooling/set up 10 of the lotion.

In applying lotion compositions of the present invention to diaper topsheets, spraying, gravure coating and extrusion coating methods are preferred. Figure 1 illustrates one such preferred method involving spraying of the coating on the diaper topsheet before the topsheet is 15 assembled with the other raw materials into a finished product. Referring to Figure 1, a nonwoven topsheet web 1 is unwound from parent topsheet roll 2 (rotating in the direction indicated by arrow 2a) and advanced to spray station 6 where one side of the web is sprayed with a hot, molten (e.g., 65°C) lotion composition. After leaving spray station 6, nonwoven topsheet 20 web 1 becomes a lotioned topsheet web indicated by 3. Lotioned topsheet web 3 is then advanced around turning roll 4 and turning roll 8, and then wound up on lotioned topsheet parent roll 10 (rotating in the direction indicated by arrow 10a).

Figure 2 illustrates an alternative preferred method involving 25 continuous or intermittent spraying of the lotion composition on to a diaper topsheet during the converting operation. Referring to Figure 2, conveyor belt 1 advances in the direction shown by the arrows on turning rolls 3 and 4 and becomes returning conveyor belt 2. Conveyor belt 1 carries nonlotioned diaper 5 to spray station 6 where topsheet patch 7 is sprayed 30 with a hot, molten (e.g., 65°C) lotion composition. After leaving spray station 6, nonlotioned diaper 5 becomes lotioned diaper 8 having a lotioned topsheet. The amount of lotion composition transferred to topsheet patch 7 is controlled by: (1) the rate at which the molten lotion composition is sprayed from spray station 6; and/or (2) the speed at which conveyor belt 1 35 travels under spray station 6.

**SPECIFIC ILLUSTRATIONS OF THE PREPARATION OF LOTIONED
DIAPER TOPSHEETS ACCORDING TO THE PRESENT INVENTION**

5 The following are specific illustrations of treating diaper topsheets with lotion compositions in accordance with the present invention:

Example 1

A. Preparation of Lotion Compositions

10 A water free lotion composition (Lotion A) is made by mixing the following melted (i.e., liquid) components together: Mineral Oil (Carnation White Mineral Oil, USP made by Witco Corp.), Cetearyl Alcohol (a mixed linear C₁₆-C₁₈ primary alcohol made by the Procter & Gamble Company under the name TA-1618); Steareth-2 (Brij 72, a C₁₈ linear alcohol ethoxylate having an average degree of 15 ethoxylation of 2, made by ICI America). The weight percentages of these components are shown in Table I below:

Table I

Component	Weight %
Mineral Oil	50
Cetearyl Alcohol	35
Steareth-2	15

B. Preparation of Lotioned Diaper Topsheet by Hot Melt Spraying

20 Lotion Composition A is placed into a heated tank operating at a temperature of 125°F. The composition is subsequently sprayed (using a Dynatec E84B1758 spray head, operating at a temperature of 165°F and an atomization pressure of 2.40 psig) onto the topsheet of a diaper in a 3.75 inch wide (diaper lateral direction) and 7 inch long (diaper longitudinal direction) area, the patch beginning 1 inch forward of the lateral centerline and extending toward the rear of the product. Add-on level = 0.006 g/in² (9.3 g/m²).

Example 2

5 The lotion composition A (prepared in accordance with the procedure in Example 1) is subsequently sprayed onto the topsheet of a diaper in a 3.75 inch wide (diaper lateral direction) stripe centered on the longitudinal centerline and extending the entire length of the product. Add-on level = 0.006 g/in² (9.3 g/m²).

Example 3

10 15 The lotion composition A (prepared in accordance with the procedure in Example 1) is subsequently sprayed onto the topsheet of a diaper in a 3.75 inch wide (diaper lateral direction) stripe centered on the longitudinal centerline and 7 inch long (diaper longitudinal direction) area, the patch beginning 1 inch forward of the lateral centerline and extending toward the rear of the product. Add-on level = 0.003 g/in² (4.65 g/m²).

Example 4A. Preparation of Lotion Compositions

20 25 A water free lotion composition (Lotion B) is made by mixing the following melted (i.e., liquid) components together: Mineral Oil (Carnation White Mineral Oil, USP made by Witco Corp.), and Cetearyl Alcohol (a mixed linear C₁₆-C₁₈ primary alcohol made by the Procter & Gamble Company under the name TA-1618). The weight percentages of these components are shown in Table II below:

Table II

Component	Weight %
Mineral Oil	65
Cetearyl Alcohol	35

B. Preparation of Lotioned Tissue by Hot Melt Spraying

Lotion Composition B is placed into a heated tank operating at a temperature of 125°F. The composition is subsequently sprayed (using a Dynatec E84B1758 spray head, operating at a temperature of 165°F and an atomization pressure of 2.40 psig) onto the topsheet of a diaper in a 3.75 inch wide (diaper lateral direction) and 7 inch long (diaper longitudinal direction) area, the patch beginning 1 inch forward of the lateral centerline and extending toward the rear of the product. Add-on level = 0.006 g/in² (9.3 g/m²).

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Example 5

A. Preparation of Lotion Composition

A water free lotion composition (Lotion C) is made by mixing the following melted (i.e., liquid) components together: White Protopet® 1S (white petrolatum made by Witco Corp.), Cetearyl Alcohol (a mixed linear C₁₆-C₁₈ primary alcohol made by the Procter & Gamble Company under the name TA-1618); Steareth-2 (Brij 762, a C₁₈ linear alcohol ethoxylate having an average degree of ethoxylation of 2 made by ICI America). The weight percentages of these components are shown in Table I below:

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A water free lotion composition (Lotion C) is made by mixing together the following melted (i.e., liquid) components in the weight percentages shown in the Table III below according to the procedure of Example 2:

Table III

Component	Weight %
WhiteProtopet® 1S	50
Cetearyl Alcohol	35
Steareth-2	15

B. Preparation of Lotioned Tissue by Hot Melt Spraying

Lotion Composition C is placed into a heated tank operating at a temperature of 125°F. The composition is subsequently sprayed (using a Dynatec E84B1758 spray head, operating at a temperature of 165°F and an atomization pressure of 2.40 psig) onto the topsheet of a diaper in a 3.75 inch wide (diaper lateral direction) and 7 inch long (diaper longitudinal direction) area, the patch beginning 1 inch forward of the lateral centerline and extending toward the rear of the product. Add-on level = 0.006 g/in² (9.3 g/m²).

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Example 6

A. Preparation of Lotion Composition

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A water free lotion composition (Lotion D) is made by mixing the following melted (i.e., liquid) components together: White Protopet® 1S (white petrolatum made by Witco Corp.); Dow Corning 556 Cosmetic Grade Fluid(a polyphenylmethylsiloxane made by the Dow Corning Corporation), An example of a particularly preferred paraffin wax is Parrafin S.P. 434 (a paraffin wax made by Strahl and Pitsch Inc.); Cetearyl Alcohol (a mixed linear C₁₆-C₁₈ primary alcohol made by the Procter & Gamble Company under the name TA-1618); PEG 2000 (a polyethylene glycol having a MW of 2000 made by Sigma-Aldrich Corp). The weight percentages of these components are shown in Table IV below:

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Table IV

Component	Weight %
WhiteProtopet® 1S	52
Polyphenylmethylsiloxane	20
Paraffin Wax	15
Cetearyl Alcohol	10
PEG 2000	3

B. Preparation of Lotioned Tissue by Hot Melt Spraying

Lotion Composition D is placed into a heated tank operating at a temperature of 150°F. The composition is subsequently sprayed (using a Dynatec E84B1758 spray head, operating at a temperature of 170°F and an atomization pressure of 2.40 psig) onto the topsheet of a diaper in a 3.75 inch wide (diaper lateral direction) and 7 inch long (diaper longitudinal direction) area, the patch beginning 1 inch forward of the lateral centerline and extending toward the rear of the product. Add-on level = 0.006 g/in² (9.3 g/m²).

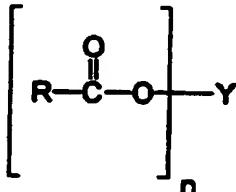
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WHAT IS CLAIMED IS:

1. **A disposable diaper characterized in that it comprises:**
 - a) **a liquid impervious backsheet;**
 - b) **a liquid pervious, hydrophilic topsheet joined to said backsheet, said topsheet having an inner surface oriented toward the interior of said diaper and an outer surface oriented toward the skin of the wearer when said diaper is being worn, wherein at least a portion of said topsheet outer surface comprises an effective amount of a lotion coating which is semi-solid or solid at 20°C and which is partially transferable to the wearer's skin, said lotion coating comprising:**
 - (i) **from 10 to 95% of a polysiloxane emollient having a plastic or fluid consistency at 20°C;**
 - (ii) **from 5 to 90%, preferably from 5 to 50% of an immobilizing agent capable of immobilizing said emollient on said outer surface of the topsheet, said immobilizing agent having a melting point of at least 35°C, preferably at least 40°C, wherein said immobilizing agent is preferably selected from polyhydroxy fatty acid esters, polyhydroxy fatty acid amides, C₁₄-C₂₂ fatty alcohols, C₁₂-C₂₂ fatty acids, C₁₂-C₂₂ fatty alcohol ethoxylates, and mixtures thereof; and**
 - c) **an absorbent core positioned between said topsheet and said backsheet.**
2. **The disposable diaper of Claim 1 wherein said emollient contains about 5% or less water and comprises a polysiloxane compound having a viscosity at 37°C of from 5 to 2,000 centistokes.**
3. **The disposable diaper of Claim 1 or 2 wherein said polysiloxane compound is a polymethylsiloxane compound, preferably said polymethylsiloxane compound is substituted with a functional group selected from phenyl, amino, alkyl, carboxyl, hydroxyl, ether, polyether, aldehyde, ketone, amide, ester, thiol groups, and mixtures thereof, more preferably said substituted polymethylsiloxane compound is a polyphenylmethylsiloxane compound.**

4. The disposable diaper of any of Claims 1-3 wherein the quantity of lotion coating on said portion of said topsheet outer surface ranges from 0.1 mg/in² to 25 mg/in², preferably from 1 mg/in² to 10 mg/in².
5. The disposable diaper of any of Claims 1-4 wherein said immobilizing agent comprises a C₁₄-C₂₂ fatty alcohol, preferably a C₁₆-C₁₈ fatty alcohol selected from cetyl alcohol, stearyl alcohol, and mixtures thereof.
6. The disposable diaper of any of Claims 1-4 wherein said immobilizing agent comprises a polyhydroxy fatty acid ester having the formula:



wherein R is a C₅-C₃₁ hydrocarbyl group; Y is a polyhydroxyhydrocarbyl moiety having a hydrocarbyl chain with at least 2 free hydroxyls directly connected to the chain; and n is at least 1, preferably said polyhydroxy fatty acid ester is selected from sorbitan esters of C₁₆-C₂₂ saturated fatty acids, glyceryl monoesters of C₁₆-C₂₂ saturated fatty acids, and sucrose esters of C₁₂-C₂₂ saturated fatty acids, most preferably said polyhydroxy fatty acid ester is selected from sorbitan palmitates, sorbitan stearates, sorbitan behenates, glyceryl monostearate, glyceryl monopalmitate, glyceryl monobehenate, sucrose mono- and di- stearate, and sucrose mono- and di- laurate.

7. The disposable diaper of any of Claims 1-4 wherein said immobilizing agent comprises a polyhydroxy fatty acid amide having the formula:



wherein R¹ is H, C₁-C₄ hydrocarbyl, 2-hydroxyethyl, 2-hydroxypropyl, methoxyethyl, methoxypropyl or a mixture thereof, preferably R¹ is N-methyl, N-ethyl, N-propyl, N-isopropyl, N-butyl, N-2-hydroxyethyl, N-

methoxypropyl or N-2-hydroxypropyl; R² is a C₅-C₃₁ hydrocarbyl group, preferably R² is straight chain C₁₁-C₁₇ alkyl or alkenyl, or mixture thereof; and Z is a polyhydroxyhydrocarbyl moiety having a linear hydrocarbyl chain with at least 3 hydroxyls directly connected to the chain, preferably glycetyl, most preferably said polyhydroxy fatty acid amide is selected from N-lauryl-N-methyl glucamide, N-lauryl-N-methoxypropyl glucamide, N-cocoyl-N-methyl glucamide, N-cocoyl-N-methoxypropyl glucamide, N-palmityl-N-methoxypropyl glucamide, N-tallowyl-N-methyl glucamide, and N-tallowyl-N-methoxypropyl glucamide.

8. The disposable diaper of any of Claims 1-7 wherein said lotion coating further comprises from 1 to 50%, preferably from 1 to 25% of a hydrophilic surfactant, said hydrophilic surfactant preferably being nonionic, said hydrophilic surfactant having an HLB value of at least 4, preferably from 4 to 20.
9. The disposable diaper of Claim 8 wherein said hydrophilic surfactant comprises an ethoxylated alcohol having an alkyl chain of from 8 to 22 carbon atoms, preferably from 11 to 22 carbon atoms, and having an average degree of ethoxylation ranging from 1 to 54, preferably from 2 to 30.
10. The disposable diaper of Claim 9 wherein said hydrophilic surfactant comprises an ethoxylated sorbitan ester of a C₁₂-C₁₈ fatty acid having an average degree of ethoxylation of from 2 to 20.

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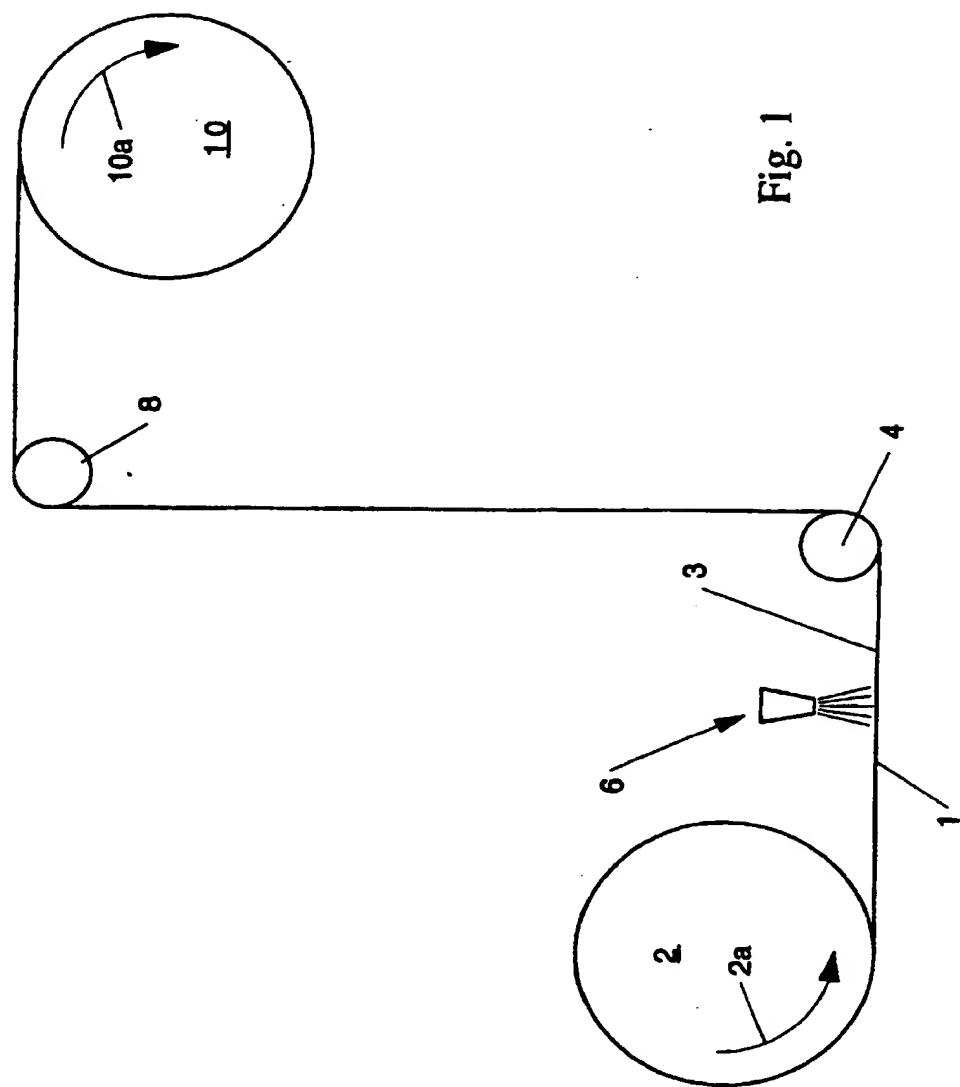


Fig. 1

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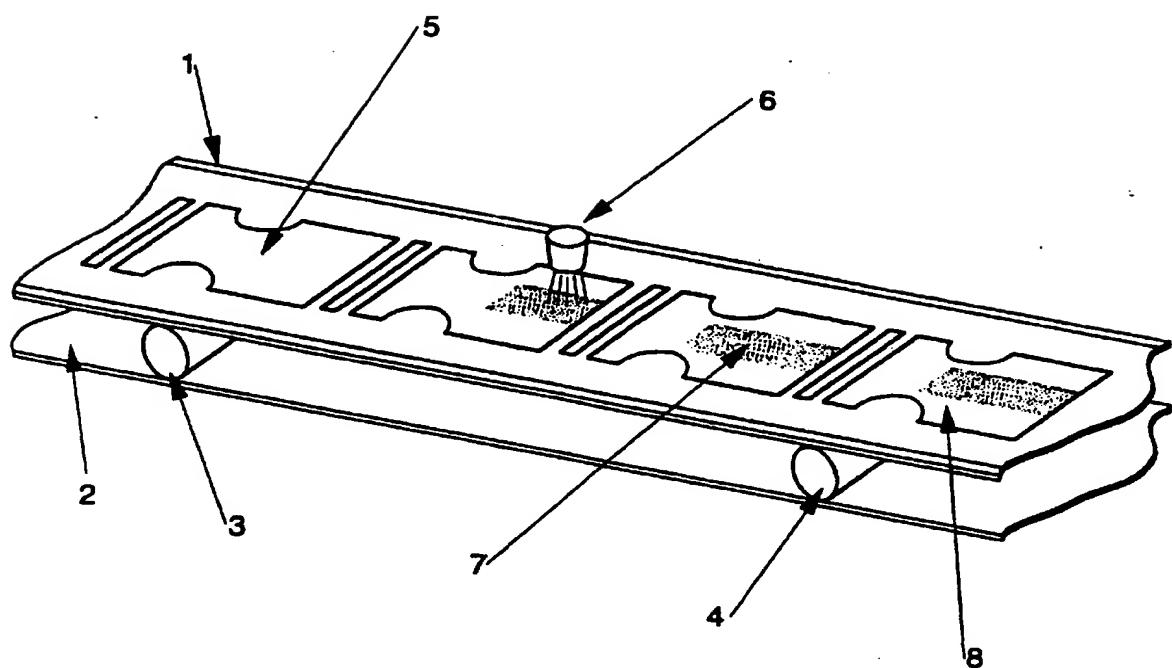


Fig. 2

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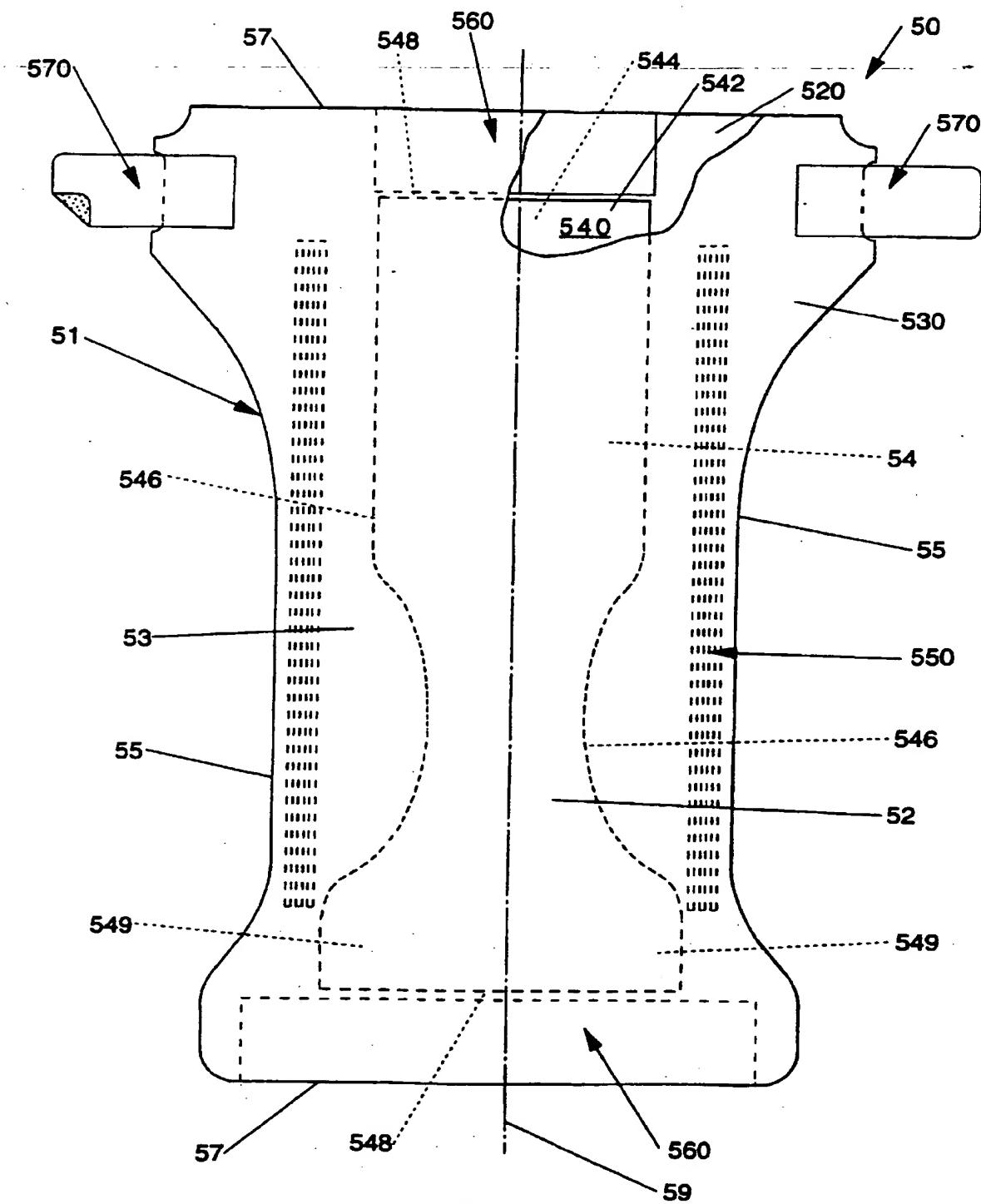


Fig. 3

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 95/13808

A. CLASSIFICATION F SUBJECT MATTER
IPC 6 A61L15/26 A61L15/34 A61L15/48

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 A61L A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO,A,94 09757 (ABBOTT LAB) 11 May 1994 see page 9, paragraph 3; claims 27-32 ---	1-10
X	DE,A,41 36 540 (AMERICAN ISRAELI PAPER MILLS) 14 May 1992 see the whole document ---	1-10
Y	US,A,3 489 148 (DUNCAN ROBERT C ET AL) 13 January 1970 cited in the application see column 2, line 25 - line 40; claims ---	1-10
A	EP,A,0 328 355 (RICHARDSON VICKS INC) 16 August 1989 ---	
Y	US,A,3 567 820 (SPERTI GEORGE S) 2 March 1971 see claims ---	1-10
	-/-	

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

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- *A* document defining the general state of the art which is not considered to be of particular relevance
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Date of the actual completion of the international search

29 March 1996

Date of mailing of the international search report

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INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 95/13808

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US,A,5 059 282 (AMPULSKI ROBERT S ET AL) 22 October 1991 ---	
A	US,A,5 013 715 (MORI SHIGERU ET AL) 7 May 1991 ---	
A	DATABASE WPI Section Ch, Week 8644 Derwent Publications Ltd., London, GB; Class A96, AN 86-287937 XP002000210 & JP,A,61 210 019 (SHISEIDO KK) , 18 September 1986 see abstract -----	

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No
PCT/US 95/13808

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
WO-A-9409757	11-05-94	US-A- 5436007		25-07-95
DE-A-4136540	14-05-92	NONE		
US-A-3489148	13-01-70	BE-A- 708264 DE-A- 1610547 FR-A- 1548038 GB-A- 1180960 NL-A- 6717354		20-06-68 08-07-71 29-11-68 11-02-70 21-06-68
EP-A-0328355	16-08-89	AU-B- 2968089 CA-A- 1329107 DE-D- 68912150 DE-T- 68912150 FI-B- 94388 JP-A- 2203929 PT-B- 89624 US-A- 5043155		10-08-89 03-05-94 24-02-94 11-08-94 31-05-95 13-08-90 29-04-94 27-08-91
US-A-3567820	02-03-71	NONE		
US-A-5059282	22-10-91	AT-T- 120820 AT-T- 132556 AT-T- 122424 AU-B- 634963 AU-B- 3636389 AU-B- 613765 AU-B- 3636489 AU-B- 634964 AU-B- 3636889 CA-A- 1328335 CA-A- 1330382 CA-A- 1328035 DE-D- 68914338 DE-T- 68914338 DE-D- 68922024 DE-T- 68922024 DE-D- 68922529 DE-T- 68922529 DE-D- 68925309		15-04-95 15-01-96 15-05-95 11-03-93 21-12-89 08-08-91 21-12-89 11-03-93 21-12-89 12-04-94 28-06-94 29-03-94 11-05-94 18-08-94 11-05-95 28-09-95 14-06-95 21-09-95 15-02-96

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No
PCT/US 95/13808

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US-A-5059282		EP-A- 0347153 EP-A- 0347154 EP-A- 0347176 EP-A- 0347177 ES-T- 2070174 ES-T- 2050802 ES-T- 2071658 JP-A- 2224626 JP-A- 2099690 JP-A- 2099691	20-12-89 20-12-89 20-12-89 20-12-89 01-06-95 01-06-94 01-07-95 06-09-90 11-04-90 11-04-90
US-A-5013715	07-05-91	JP-C- 1786523 JP-B- 4076389 JP-A- 63235366	10-09-93 03-12-92 30-09-88